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5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21	FOR THE CENTRAL DI EASTER RONALD COUWENHOVEN, Plaintiff, vs. ABBOTT LABORATORIES, INC., ABBVIE, INC., ACTAVIS, INC.,	S DISTRICT COURT ISTRICT OF CALIFORNIA IN DIVISION Case No. Sing-cv-lean Jeb-Dib COMPLAINT AND DEMAND FOR JURY TRIAL					
20	vs. ABBOTT LABORATORIES, INC.,	JURY IRIAL					
23	JOHN DOE DRUG COMPANY						
24	DEFENDANTS (1-50), JANE DOE						
25	DRUG DISTRIBUTOR DEFENDANTS (1-50), JIM DOE						
26	DOE HEALTH CARE PROVIDERS						
27	(1-50), and JILL DOE (1-50),						
28	Defendants.						
	COMPLAINT FOR DAMAGES						

Plaintiff RONALD COUWENHOVEN ("Plaintiff"), files this Complaint against the Defendants, ABBOTT LABORATORIES, INC., ABBVIE, INC., ACTAVIS, INC., and DOES (all defendants collectively hereinafter referred to as the "Defendants") alleging as follows:

NATURE OF ACTION

- 1. This is an action for damages suffered by Plaintiff as a direct and proximate result of Defendants' wrongful conduct in connection with the development, design, testing, labeling, packaging, promoting, advertising, marketing, distribution, and selling of Defendants' prescription medication Androgel and Androderm.
- 2. This case involves the prescription drugs Androgel and Androderm ("testosterone"), which are manufactured, sold, distributed and promoted by Defendants as testosterone replacement therapies.
- 3. Defendants misrepresented that testosterone is a safe and effective treatment for hypogonadism or "low testosterone," when in fact these drugs cause serious medical problems, including life threatening cardiac events, strokes, and thrombolytic events.
- 4. Defendants engaged in aggressive, award-winning direct-to-consumer and physician marketing and advertising campaigns for testosterone. Further, Defendants engaged in an aggressive unbranded "disease awareness" campaign to alert men that they might be suffering from "low T."
- 5. As a result, diagnoses of Low T and prescriptions for testosterone replacement therapies have increased exponentially. For example:
 - a. Defendants ABBOTT LABORATORIES, INC. ("ABBOTT")
 and ABBVIE, INC.'s sales of AndroGel have increased to over
 \$1.37 billion per year; and

- b. Defendant ACTAVIS, INC.'s sales of Androderm have increased to over \$69 million per year;
- 6. However, consumers of testosterone were misled as to the drug's safety and efficacy, and as a result have suffered injuries including life-threatening cardiac events, strokes, and thrombolytic events.

THE PARTIES

- 7. Plaintiff RONALD COUWENHOVEN ("COUWENHOVEN") is a citizen of the State of California, and a resident of Mira Loma, California, county of Riverside.
- 8. Upon information and belief, Defendant ABBOTT, a manufacturer of AndroGel, is a corporation organized and existing under the laws of Illinois and maintains its principal place of business at 100 Abbott Park Road, Abbott Park, Illinois 60064. ABBOTT has conducted business and derived substantial revenue from within the State of California.
- 9. Upon information and belief, Defendant ABBVIE, INC., a manufacturer of AndroGel, is a corporation organized and existing under the laws of Delaware with its principal place of business at 1 North Waukegan Road, North Chicago, Illinois 60064. ABBVIE, INC. has conducted business and derived substantial revenue from within the State of California.
- developed AndroGel and sought FDA approval in 1999. Before the drug was approved by the FDA in 2000, Solvay Pharmaceuticals Inc. acquired Unimed Pharmaceuticals, Inc. and subsequently brought AndroGel to market. In 2010, Defendant ABBOTT acquired Solvay's pharmaceutical division, which included AndroGel. Then, in 2013, ABOTT created AbbVie, a company composed of Abbott's former proprietary pharmaceutical business, which included AndroGel.
- 11. Upon information and belief, Defendant ACTAVIS, INC., a manufacturer of Androderm, is a corporation organized and existing under the laws

- 12. By way of background, TheraTech, Inc. originally developed Androderm, which was approved by the FDA on 9/29/1995. Watson acquired TheraTech in January 1999 and continued to manufacture and distribute Androderm. Watson then acquired Actavis on October 31, 2012, and subsequently changed its corporate name to Actavis, Inc. on January 23, 2013.
- 13. Defendant John Doe Manufacturer Defendants are defendants who are or have been involved in the manufacture, distribution, marketing, sale and labeling of testosterone products but are not yet known by Plaintiff(s).
- 14. At all times relevant herein, Defendants tested, studied, researched, designed, formulated, manufactured, inspected, labeled, packaged, promoted, advertised, marketed, distributed, and sold the prescription drugs Androgel and Androderm in interstate commerce and throughout the State of California. _At all times relevant herein, Defendants were registered to do business in the State of California.

JURISDICTION AND VENUE

- 15. This Court has jurisdiction pursuant to 28 U.S.C. § 1332(a) because Plaintiff and Defendants are citizens of different States and the amount in controversy exceeds \$75,000 exclusive of interest and costs.
- 16. Venue in this action properly lies in this judicial district pursuant to 28 U.S.C. § 1391(a), as a substantial number of the events, actions or omissions giving rise to Plaintiff's claims occurred in this district. At all times material hereto, Defendants conducted substantial business in this district.

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TESTOSTERONE THERAPY AND ITS SIDE EFFECTS

17. Hypogonadism is a specific condition of the sex glands that may involve the diminished production or nonproduction of testosterone in males.

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- 18. Testosterone is a primary androgenic hormone responsible for normal growth, development of the male sex organs, and maintenance of secondary sex characteristics.
- 19. The hormone plays a role in sperm production, fat distribution, maintenance of muscle strength and mass, and sex drive.
- 20. In men, testosterone levels normally begin a gradual decline after the age of thirty.
- 21. The average testosterone levels for most men range from 300 to 1,000 nanograms per deciliter of blood. Testosterone levels can fluctuate greatly depending on many factors, including sleep, time of day, and medication. Resultantly, many men who fall into the hypogonadal range one day will have normal testosterone levels the next.

Androgel

- 22. The Food and Drug Administration approved AndroGel 1% on February 28, 2000, and then approved AndroGel 1.62% on April 29, 2011. After FDA approval, AndroGel was widely advertised and marketed as a safe and effective testosterone replacement therapy.
- 23. AndroGel is a hydroalcoholic gel containing testosterone thatis applied to the shoulders and upper arms, and enters the body through transdermal absorption.
- 24. AndroGel may produce undesirable side effects to patients who use the drug, including, but not limited to, myocardial infarction, stroke, and death.
- 25. In some patient populations, AndroGel use may increase the incidence of myocardial infarctions and death by more than 500%.

COMPLAINT FOR DAMAGES

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Androderm

- 26. The Food and Drug Administration approved Androderm on September 29, 1995. After FDA approval, Androderm was widely advertised and marketed as a safe and effective testosterone replacement therapy.
- 27. Androderm is a transdermal patch containing testosterone andis applied to the back, abdomen, upper arms and thighs.
- 28. Androderm may produce undesirable side effects to patients who use the drug, including, but not limited to, myocardial infarction, stroke, and death.
- 29. In some patient populations, Androderm use may increase the incidence of myocardial infarctions and death by more than 500%.
- 30. In 2010, a New England Journal of Medicine Study entitled "Adverse Events Associated with Testosterone Administration" was discontinued after an exceedingly high number of men suffered adverse events.
- 31. In November 2013, a JAMA study was released entitled "Association of Testosterone Therapy with Mortality, Myocardial Infarction, and Stroke in Men with Low Testosterone Levels," which indicated that testosterone therapy raised the risk of death, heart attack and stroke by approximately 30%.
- 32. On January 29, 2014, a study was released in PLOS ONE entitled "Increased Risk of Non-Fatal Myocardial Infarction Following Testosterone Therapy Prescription in Men," which indicated that testosterone use doubled the risk of heart attacks in men over sixty-five years old and men younger than sixty-five with a previous diagnosis of heart disease.
- 33. There have been a number of studies concluding that testosterone therapy causes a sudden increase in hematocrit, hemoglobin and estradiol, and associating its use with an increased risk of heart attacks and strokes.
- 34. In addition to the above, Defendants' testosterone product has been linked to several severe and life changing medical disorders in the user of the product and in those who come into physical contact with the user or the user's

COMPLAINT FOR DAMAGES

 unwashed clothes. Patients taking an aforementioned testosterone product may experience enlarged prostates and increased serum prostate-specific antigen levels.

- 35. Secondary exposure to testosterone can cause side effects in others. In 2009, the FDA issued a black box warning for testosterone prescriptions, advising patients of reported virilization in children who were secondarily exposed to the gel. Testosterone may also cause physical changes in women exposed to the drug and cause fetal damage with pregnant women who come into secondary contact with testosterone.
- 36. Defendants' marketing strategy has been to aggressively market and sell their products by misleading potential users about the prevalence and symptoms of low testosterone and by failing to protect users from serious dangers that Defendants knew or should have known result from use of its products.
- 37. Defendants successfully marketed testosterone by undertaking a "disease awareness" marketing campaigns. These campaigns sought to create a consumer perception that low testosterone is prevalent among U.S. men and that symptoms previously associated with other physical and mental conditions, such as aging, stress, depression, and lethargy were actually attributable to "Low-T."
- 38. Defendant coordinated massive advertising campaigns designed to convince men that they suffered from low testosterone. Defendant orchestrated national disease awareness media blitzes that purported to educate male consumers about the signs of low testosterone. The marketing campaigns included promotional literature placed in healthcare providers' offices and distributed to potential testosterone users, and online media.
- 39. The advertisements suggest that various symptoms often associated with other conditions may be caused by low testosterone and encourage men to discuss testosterone replacement therapy with their doctors if they experienced any of the "symptoms" of low testosterone. These "symptoms" include listlessness,

 increased body fat, and moodiness—all general symptoms that are often a result of aging, weight gain, or lifestyle, rather than low testosterone.

- 40. Defendants' advertising programs sought to create the image and belief by consumers and their physicians that the use of testosterone was a safe method of alleviating their symptoms, had few side effects and would not interfere with their daily lives, despite that Defendants knew or should have known these to be false and without any support from their own studies or widely accepted medical literature.
- 41. Defendants purposefully downplayed, understated and outright ignored the health hazards and risks associated with using testosterone.

 Defendants deceived potential testosterone users by relaying positive information through the press, including testimonials from retired professional athletes, and manipulating hypogonadism statistics to suggest widespread disease prevalence, while downplaying known adverse and serious health effects.
- 42. Defendants concealed material relevant information from potential testosterone users and minimized user and prescriber concern regarding the safety of testosterone replacement therapy.
- 43. In particular, Defendants fail to mention any potential cardiac or stroke side effects in their commercials, online and print advertisements, and falsely represent that Defendants adequately tested testosterone for all likely side effects.
- 44. As a result of Defendants' advertising and marketing, and representations about their products, men in the United States pervasively seek out prescriptions for testosterone. If Plaintiff in this action had known the risks and dangers associated with testosterone, Plaintiff would not have taken testosterone and consequently would not have been subject to its serious side effects.

- 45. Defendants also sought to convince primary care physicians that low testosterone levels are widely under-diagnosed, and that conditions associated with normal aging could be caused by low testosterone levels.
- 46. A study published in the Journal of the American Medical Association ("JAMA") in August 2013 entitled "Trends in Androgen Prescribing in the United States, 2001-2011" indicated that many men who get testosterone prescriptions have no evidence of hypogonadism. For example, one third of men prescribed testosterone had a diagnosis of fatigue, and one quarter of men did not even have their testosterone levels tested before they received a testosterone prescription.
- While running disease awareness campaigns, Defendants promoted their testosterone product as an easy to use topical testosterone replacement therapy. Defendants contrast their products' at-home topical application with less convenient prescription testosterone injections, which require frequent doctor visits.
- 48. Defendants convinced millions of men to discuss testosterone replacement therapy with their doctors, and consumers and their physicians relied on Defendants' promises of safety. Although prescription testosterone replacement therapy has been available for years, it was not until Defendants' massive marketing campaign that millions of men, who were never been prescribed testosterone, flocked to their doctors and pharmacies.
- 49. What consumers received, however, were not safe drugs, but products that cause life-threatening problems, including strokes, heart attacks, and death.
- 50. Defendants successfully created a robust and previously nonexistent market for their drugs. Defendants spent millions of dollars promoting their products. Defendants also spent millions on their unbranded marketing including commercials and websites recommending that men have regular checkups with their physicians and five regular tests done: tests for cholesterol, blood pressure, blood sugar, prostate-specific antigen, and testosterone.

- 52. The Defendants' marketing program sought to create the image and belief by consumers and physicians that low testosterone affected a large number of men in the United States and that the use of testosterone is safe for human use, despite that Defendants knew these to be false and without any support from their own studies or widely accepted medical literature.
- 53. Defendants engaged in aggressive, award-winning direct-to-consumer and physician marketing and advertising campaigns for testosterone. Further, Defendants engaged in an aggressive unbranded "disease awareness" campaign to alert men that they might be suffering from "low T."

SPECIFIC FACTUAL ALLEGATIONS

- 54. In or about 2007, Plaintiff RONALD COUWENHOVEN was sixty two years of age when he was prescribed and began Androgel and/or Androderm for symptoms he attributed to low testosterone after viewing Defendants' advertisements. Plaintiff started taking Androgel and/or Androderm in or about September 2007 and stopped taking it in in or about 2014.
- 55. Neither Plaintiff, nor his physician, received an adequate warning from Defendants about the risk of persistent and/or permanent injury after discontinuation of treatment.
- 56. Plaintiff was very healthy and had no history of blood clots prior to taking testosterone. In keeping with his healthy and proactive lifestyle, Plaintiff agreed to initiate testosterone treatment. He relied on claims made by Defendants

that testosterone had been clinically shown to safely and effectively raise testosterone levels.

57. Plaintiff was diagnosed with Deep Vein Thrombosis on or about April 5, 2012. As a result, Plaintiff was prescribed blood thinners, etc.

CLAIMS FOR RELIEF FIRST CLAIM FOR RELIEF

Strict Liability - Failure to Warn

- 58. Plaintiff incorporates by reference each and every paragraph of this Complaint as if fully set forth herein and further alleges as follows:
- Defendants were defective due to inadequate warnings or instructions because Defendants knew or should have known that the products created significant risks of serious bodily harm to consumers, and they failed to adequately warn consumers and/or their health care providers of such risks. The testosterone products manufactured and/or supplied by Defendants were defective due to inadequate post-marketing warnings or instructions because, after Defendants knew or should have known of the risk of serious bodily harm from the use of testosterone, Defendants failed to provide an adequate warning to consumers and/or their health care providers of the product.
- 60. As a direct and proximate result of Plaintiff's reasonably anticipated use of testosterone as manufactured, designed, sold, supplied, marketed and/or introduced into the stream of commerce by Defendants, Plaintiff suffered serious injury, harm, damages, economic and non-economic loss and will continue to suffer such harm, damages and losses in the future.

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SECOND CLAIM FOR RELIEF

Negligence

- 61. Plaintiff incorporates by reference each and every paragraph of this Complaint as if fully set forth herein and further alleges as follows: At all times herein mentioned, Defendants had a duty to properly manufacture, design, formulate, compound, test, produce, process, assemble, inspect, research, distribute, market, label, package, distribute, prepare for use, sell, prescribe and adequately warn of the risks and dangers of testosterone.
- 62. At all times herein mentioned, Defendants negligently and carelessly manufactured, designed, formulated, distributed, compounded, produced, processed, assembled, inspected, distributed, marketed, labeled, packaged, prepared for use and sold testosterone and failed to adequately test and warn of the risks and dangers of testosterone.
- 63. Despite the fact that Defendants knew or should have known that testosterone caused unreasonable, dangerous side effects, Defendants continued to market testosterone to consumers, including Plaintiff, when there were safer alternative methods of treating loss of energy, libido erectile dysfunction, depression, loss of muscle mass and other conditions that the testosterone advertising claims are caused by low testosterone.
- 64. Defendants knew or should have known that consumers such as Plaintiff would foreseeably suffer injury as a result of Defendants' failure to exercise ordinary care as described above.
- 65. As a direct and proximate cause of Defendants' negligence, Plaintiff suffered serious injury, harm, damages, economic and non-economic loss and will continue to suffer such harm, damages and losses in the future.

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THIRD CLAIM FOR RELIEF

Breach of Implied Warranty

- 66. Plaintiff incorporates by reference each and every paragraph of this Complaint as if fully set forth herein and further alleges as follows:
- 67. Prior to the time that the aforementioned products were used by Plaintiff, Defendants impliedly warranted to Plaintiff and Plaintiff's agents and physicians that testosterone was of merchantable quality and safe and fit for the use for which it was intended.
- 68. Plaintiff was and is unskilled in the research, design and manufacture of the products and reasonably relied entirely on the skill, judgment and implied warranty of the Defendants in using testosterone.
- 69. Testosterone was neither safe for its intended use nor of merchantable quality, as warranted by Defendants, in that testosterone has dangerous propensities and will cause severe injuries to users when used as intended.
- 70. As a result of the abovementioned breach of implied warranties by Defendants, Plaintiff suffered serious injury, harm, damages, economic and non-economic loss and will continue to suffer such harm, damages and losses in the future.

FOURTH CLAIM FOR RELIEF

Breach of Express Warranty

- 71. Plaintiff incorporates by reference each and every paragraph of this Complaint as if fully set forth herein and further alleges as follows:
- 72. At all times mentioned, Defendants expressly represented and warranted to Plaintiff and Plaintiff's agents and physicians, by and through statements made by Defendants or their authorized agents or sales representatives, orally and in publications, package inserts and other written materials intended for physicians, medical patients and the general public, that testosterone is safe,

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- 73. In utilizing testosterone, Plaintiff relied on the skill, judgment, representations, and foregoing express warranties of Defendants. These warranties and representations were false in that testosterone is unsafe and unfit for its intended uses.
- 74. As a result of the abovementioned breach of express warranties by Defendants, Plaintiff suffered serious injury, harm, damages, economic and non-economic loss and will continue to suffer such harm, damages and losses in the future.

FIFTH CLAIM FOR RELIEF

Fraud

- 75. Plaintiff incorporates by reference each and every paragraph of this Complaint as if fully set forth herein and further alleges as follows:
- 76. Defendants, from the time they first tested, studied, researched, evaluated, endorsed, manufactured, marketed and distributed testosterone, and up to the present, willfully deceived Plaintiff, Plaintiff's physicians and the general public, by concealing from them the true facts concerning testosterone, which the Defendants had a duty to disclose.
- 77. At all times herein mentioned, Defendants conducted a sales and marketing campaign to promote the sale of testosterone and willfully deceived Plaintiff, Plaintiff's physicians and the general public as to the benefits, health risks and consequences of using testosterone. Defendants knew of the foregoing, that testosterone is not safe, fit and effective for human consumption, that using testosterone is hazardous to health, and that testosterone has a serious propensity to cause serious injuries to its users including, but not limited to, the injuries Plaintiff suffered.

78. Defendants concealed and suppressed the true facts concerning testosterone with the intent to defraud Plaintiff, in that Defendants knew that Plaintiff physicians would not prescribe testosterone, and Plaintiff would not have used testosterone, if they were aware of the true facts concerning its dangers.

79. As a result of Defendants' fraudulent and deceitful conduct, Plaintiff suffered serious injury, harm, damages, economic and non-economic loss and will continue to suffer such harm, damages and losses in the future.

SIXTH CLAIM FOR RELIEF

Negligent Misrepresentation

- 80. Plaintiff incorporates by reference each and every paragraph of this Complaint as if fully set forth herein and further alleges as follows:
- 81. From the time testosterone was first tested, studied, researched, evaluated, endorsed, manufactured, marketed and distributed, and up to the present, Defendants made misrepresentations to Plaintiff, Plaintiff's physicians and the general public including, but not limited to, the misrepresentation that testosterone was safe, fit and effective for human consumption. At all times mentioned, Defendants conducted sales and marketing campaigns to promote the sale of testosterone and willfully deceived Plaintiff, Plaintiff's physicians and the general public as to the health risks and consequences of the use of the abovementioned products.
- 82. The Defendants made the foregoing representation without any reasonable ground for believing them to be true. These representations were made directly by Defendants, by sales representatives and other authorized agents of Defendants, and in publications and other written materials directed to physicians, medical patients and the public.
- 83. The representations by the Defendants were in fact false, in that testosterone is not safe, fit and effective for human consumption, using testosterone

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is hazardous to health, and testosterone has a serious propensity to cause serious injuries to users, including but not limited to the injuries suffered by Plaintiff.

- The foregoing representations by Defendants were made with the 84 intention of inducing reliance and the prescription, purchase and use of testosterone.
- In reliance of the misrepresentations by the Defendants, Plaintiff was 85. induced to purchase and use testosterone. If Plaintiff had known of the true facts and the facts concealed by the Defendants, Plaintiff would not have used testosterone. The reliance of Plaintiff upon Defendants' misrepresentations was justified because such misrepresentations were made and conducted by individuals and entities that were in a position to know the true facts.
- As a result of the foregoing negligent misrepresentations by 86. Defendants, Plaintiff suffered serious injury, harm, damages, economic and noneconomic loss and will continue to suffer such harm, damages and losses in the future.

SEVENTH CLAIM FOR RELIEF

Fraudulent Concealment

- Plaintiff incorporates by reference each and every paragraph of this 87. Complaint as if fully set forth herein and further alleges as follows:
- Any applicable statutes of limitations have been tolled by the knowing and active concealment and denial of material facts known by each Defendant when it had a duty to disclose those facts. Each Defendant has kept Plaintiff ignorant of vital information essential to his pursuit of these claims, without any fault or lack of diligence on Plaintiff's part, for the purpose of obtaining delay on Plaintiff's part in filing a complaint on the causes of action. Each Defendants' fraudulent concealment did result in such delay. Plaintiff could not reasonably have discovered these claims until shortly before filing his original complaint.

89. Each Defendant was under a continuing duty to disclose the true character, quality, and nature of its drug that Plaintiff utilized, but instead concealed them. As a result, each Defendant is estopped from relying on any statute of limitations defense.

EIGHTH CLAIM FOR RELIEF

Violation of Unfair and Deceptive Trade Practices Act

- 90. Plaintiff incorporates by reference each and every paragraph of this Complaint as if fully set forth herein and further alleges as follows:
- 91. Defendants have a statutory duty to refrain from unfair or deceptive acts or trade practices in the design, development, manufacture, promotion, and sale of testosterone.
- 92. Had the Defendants not engaged in the deceptive conduct described herein, Plaintiff would not have purchased and/or paid for testosterone, and would not have incurred related medical costs. Specifically, Plaintiff, his physician, and Plaintiff's physician's staff were misled by the deceptive conduct described herein.
- 93. Defendants' deceptive, unconscionable, and/or fraudulent representations and material omissions to patients, physicians and consumers, including Plaintiff, constituted unfair and deceptive acts and trade practices in violation of the state consumer protection statute listed below.
- 94. Defendants engaged in wrongful conduct while at the same time obtaining, under false pretenses, substantial sums of money from Plaintiff for testosterone that they would not have paid had Defendants not engaged in unfair and deceptive conduct.
- 95. Plaintiff was injured by the cumulative and indivisible nature of Defendants' conduct. The cumulative effect of Defendants' conduct directed at patients, physicians and consumers was to create a demand for and sell testosterone. Each aspect of Defendants' conduct combined to artificially create sales of testosterone.

- 96. The medical community relied upon Defendants' misrepresentations and omissions in determining to use testosterone.
- 97. By reason of the unlawful acts engaged in by Defendants, Plaintiff has suffered ascertainable loss and damages.
- 98. As a direct and proximate result of Defendants' wrongful conduct, Plaintiff was damaged by paying in whole or in part for testosterone.
- 99. As a direct and proximate result of Defendants' violations of unfair trade practice acts, Plaintiff has sustained economic losses and other damages for which he is entitled to statutory and compensatory damages, and declaratory relief, in an amount to be proven at trial.

NINTH CLAIM FOR RELIEF

Negligent Infliction of Emotional Distress

- 100. Plaintiff incorporates by reference each and every paragraph of this Complaint as if fully set forth herein and further alleges as follows:
- 101. Defendants carelessly and negligently manufactured, marketed, and sold testosterone to Plaintiff, carelessly and negligently concealed defects from Plaintiff, and carelessly and negligently misrepresented the quality and safety of testosterone. Defendants should have realized that such conduct involved an unreasonable risk of causing emotional distress to reasonable persons, that might, in turn, result in illness or bodily harm.
- 102. Defendants owed a duty to treating physicians and Plaintiff to accurately and truthfully represent the risks of testosterone. Defendants breached that duty by misrepresenting and/or failing to adequately warn of the risks of testosterone effects of which Defendants knew or in the exercise of diligence should have known to the treating physicians and Plaintiffs.
- 103. As a direct and proximate result of Defendants' wrongful conduct and breach of duty, Plaintiff has sustained and will continue to sustain severe emotional

distress either due to physical injury or a rational fear of physical injury and is entitled to recovery of damages in an amount to be proven at trial.

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NINTH CLAIM FOR RELIEF

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Loss of Consortium/Per Quod Claim

104. Plaintiff incorporates by reference each and every paragraph of this

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Complaint as if fully set forth herein and further alleges as follows:

105. By reason of the foregoing, Plaintiff's spouse has necessarily paid and

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has become liable to pay for medical aid, treatment, attendance and medications, and will necessarily incur further expenses of a similar nature in the future.

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106. By reason of the foregoing, Plaintiff's spouse has been caused presently and in the future the loss of her husband's companionship, services and society.

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PRAYER FOR RELIEF

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WHEREFORE, Plaintiff respectfully prays for relief as follows:

17 18 (a) Compensatory damages, in excess of the amount required for federal diversity jurisdiction, and in an amount to fully compensate Plaintiff for all of his injuries and damages, both past and present;

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(b) Special damages, in excess of the amount required for federal diversity jurisdiction and in an amount to fully compensate Plaintiff for all of his injuries and damages, both past and present, including but not limited to, past and future medical expenses, costs for past and future rehabilitation and/or home health care, lost income, permanent disability, including permanent instability and loss of

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balance, and pain and suffering.

(c) Punitive and/or exemplary damages for the wanton, willful, fraudulent, reckless acts of Defendant who demonstrated a complete disregard and reckless indifference for the safety and welfare of the general public and to

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COMPLAINT FOR DAMAGES

Plaintiff in an amount sufficient to punish Defendant and deter future similar 1 conduct 2 Double or triple damages as allowed by law; (d) 3 Attorneys' fees, expenses, and costs of this action; 4 (e) Pre-judgment and post-judgment interest in the maximum amount (f) 5 allowed by law; and 6 Such further relief as this Court deems necessary, just, and proper. 7 (g) 8 9 Dated: April 4, 2014 THE LANIER LAW FIRM 10 11 12 Lee A. Cirsch 13 Michael Akselrud 2049 Century Park East, Suite 1940 Los Angeles, California 90067 Phone: (310) 277-5100 Fax: (310) 277-5103 lee.cirsch@lanierlawfirm.com 14 15 16 michael.akselrud@lanierlawfirm.com 17 18 W. Mark Lanier Catherine Heacox 19 126 East 56th Street, 6th Floor New York, NY 10022 Phone: (212) 421-2800 20 21 Fax: (212) 421-2878 wml@lanierlawfirm.com 22 catherine.heacox@lanierlawfirm.com 23 Attorneys for Plaintiff 24 RONALD COUWENHOVEN 25 26 27 28

JURY DEMAND Plaintiff RONALD COUWENHOVEN hereby demands a trial by jury. THE LANIER LAW FIRM ee A. Cirsch Michael Akselrud 2049 Century Park East, Suite 1940 Los Angeles, California 90067 Phone: (310) 277-5100 Fax: (310) 277-5103 lee.cirsch@lanierlawfirm.com michael.akselrud@lanierlawfirm.com W. Mark Lanier Catherine Heacox 126 East 56th Street, 6th Floor New York, NY 10022 Phone: (212) 421-2800 Fax: (212) 421-2878 wml@lanierlawfirm.com catherine.heacox@lanierlawfirm.com Attorneys for Plaintiff RONALD COUWENHOVEN

UNITED STATES DISTRICT COURT, CENTRAL DISTRICT OF CALIFORNIA **CIVIL COVER SHEET**

I. (a) PLAINTIFFS (Che	ck box if you are repres	senting yourself [)	DEFENDANTS	(Check box if you are rep	resenting yourself 🔲)			
RONALD COUWENHOVEN, ar	ı individual.		ABBOTT LABORATOR	ABBOTT LABORATORIES, INC, et al.				
(b) County of Residence	of First Listed Plaint	tiff Riverside	County of Resider	nce of First Listed Defen	dant			
(EXCEPT IN U.S. PLAINTIFF CASI	ES)		(IN U.S. PLAINTIFF CAS	ES ONLY)				
(c) Attorneys (Firm Name representing yourself, pro Lee A. Cirsch (SBN 227668), N THE LANIER LAW FIRM, PC, 20 Phone: (310) 277-5100; Fax: (vide the same informa Aichael A. Akselrud (SBN 2 049 Century Park East, Sui 310) 277-5103	tion. !85033) ite 1940		ame, Address and Telephone elf, provide the same inforr				
lee.cirsch@lanierlawfirm.com		1	U CITIZENCI UD OF DD	INCIDAL DARTICE C	ity Casa Only			
II. BASIS OF JURISDIC	TION (Place an X in or	ne box only.)	(Place an X in one box	INCIPAL PARTIES-For Di x for plaintiff and one for de	efendant)			
1. U.S. Government	3. Federal Qu			ren of This State PTF DEF Incorporated or Principal Place of Business in this State PTF DEF 4 × 4				
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2. U.S. Government	Tal A Diversity (I	ndicate Citizenship C	itizen or Subject of a	of Business in Ar				
Defendant	of Parties in It	tem III)	oreign Country] 3 📋 3 Foreign Nation	<u> 6 6 </u>			
V		3. Remanded from Appellate Court	Reopened Dis	nsferred from Another Strict (Specify)	Multi- District igation			
V. REQUESTED IN COM	иPLAINT: JURY DE	MAND: X Yes	No (Check "Yes" or	nly if demanded in comp	olaint.)			
CLASS ACTION under	F.R.Cv.P. 23:	∕es ⊠No	MONEY DEMA	NDED IN COMPLAINT:	\$			
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VII. NATURE OF SUIT			INMUCRATION	DDICOMED DETITIONS	DRODERTY RIGHTS			
OTHER STATUTES	CONTRACT	REAL PROPERTY CONT.	The state of the s	PRISONER PETITIONS	PROPERTY RIGHTS			
OTHER STATUTES 375 False Claims Act	CONTRACT 110 Insurance	REAL PROPERTY CONT. 240 Torts to Land	IMMIGRATION 462 Naturalization Application	Habeas Corpus:	820 Copyrights			
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CIVIL COVER SHEET

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UNITED STATES DISTRICT COURT, CENTRAL DISTRICT OF CALIFORNIA CIVIL COVER SHEET

VIII. VENUE: Your answers to the questions below will determine the division of the Court to which this case will most likely be initially assigned. This initial assignment is subject to change, in accordance with the Court's General Orders, upon review by the Court of your Complaint or Notice of Removal.

Question A: Was this case removed from state court?			STATE CASE WAS PENDING IN THE COUNTY OF:				INITIAL DIVISION IN CACD IS:		
Yes X No			os Angeles				Western		
If "no, " go to Question B. If "yes," check the box to the right that applies, enter the corresponding division in response to			Ventura, Santa Barbara, or San Luis Obispo				Western		
			range					Southern	
Question D, below, and skip to Sectior	ı IX.	Ri	verside or San Bernardino				Eastern		
Question B: Is the United States, or one of its agencies or employees, a party to this action? Yes No If "no, " go to Question C. If "yes," check the box to the right that applies, enter the corresponding division in response to Question D, below, and skip to Section IX.		A PLAINTIFF? Then check the box below for the county in which the majority of DEFENDANTS reside. Los Angeles Ventura, Santa Barbara, or San Luis Obispo Orange		unty in 5 reside,	A DEFENDANT? Then check the box below for the county in which the majority of PLAINTIFFS reside. Los Angeles Ventura, Santa Barbara, or San Luis Obispo Orange		INITIAL DIVISION IN CACD IS: Western Western Southern		
		Riverside or San Bernardino			Riverside or San Bernardino		Eastern		
			Other		Other		Western		
Question C: Location of plaintiffs, defendants, and claims? (Make only one selection per row)	Los Ar Cou	ngeles	B. Ventura, Santa Barbara, or San Luis Obispo Counties	C: Orange Co	ounty	D. Riverside or San Bernardino Counties		E. de the Central et of California	F. Othe
Indicate the location in which a majority of plaintiffs reside:						×			
Indicate the location in which a majority of defendants reside:							X		
Indicate the location in which a majority of claims arose:						X	X		
		198							Line
C.1. Is either of the following true? If so, check the one that applies: 2 or more answers in Column C only 1 answer in Column C and no answers in Column D Your case will initially be assigned to the SOUTHERN DIVISION. Enter "Southern" in response to Question D, below. If none applies, answer question C2 to the right.				C.2. Is either of the following true? If so, check the one that applies: 2 or more answers in Column D only 1 answer in Column D and no answers in Column C Your case will initially be assigned to the EASTERN DIVISION. Enter "Eastern" in response to Question D, below. If none applies, go to the box below.					
			Your case will WES Enter "Western" in	TERN DIVISI	ON.				
						INITIAL DIVIS	ION IN CA	CD	50.7
Question D: Initial Division?				Eastern Division					

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UNITED STATES DISTRICT COURT, CENTRAL DISTRICT OF CALIFORNIA CIVIL COVER SHEET

IX(a). IDENTICAL CA	SES: Has this act	ion been previously filed in this court a nd dismissed, remanded or closed?	⋈ NO	YES
If yes, list case num	nber(s):			
IX(b). RELATED CASI	ES : Have any case	es been previously filed in this court that are related to the present case?	⊠ NO	YES
If yes, list case num	nber(s):			
Civil cases are deemed	I related if a previo	usly filed case and the present case:		
(Check all boxes that ap	ply) A. Arise f	rom the same or closely related transactions, happenings, or events; or		
	B. Call fo	r determination of the same or substantially related or similar questions of law and fac	t; or	
	C. For ot	her reasons would entail substantial duplication of labor if heard by different judges; o	r	
	D. Involv	ve the same patent, trademark or copyright <u>, and</u> one of the factors identified above in a	a, b or c also is pre	esent.
Notice to Counsel/Partie	S: The CV-71 (JS-44) by law. This form, ap the Court for the pu	Civil Cover Sheet and the information contained herein neither replace nor supplemer proved by the Judicial Conference of the United States in September 1974, is required troose of statistics, venue and initiating the civil docket sheet. (For more detailed instru	pursuant to Loca	I RUJE 3-1 IS NOT THEG
Nature of Suit Code		Substantive Statement of Cause of Action		
861	HIA	All claims for health insurance benefits (Medicare) under Title 18, Part A, of the Social include claims by hospitals, skilled nursing facilities, etc., for certification as provider (42 U.S.C. 1935FF(b))	al Security Act, as rs of services und	amended. Also, er the program.
862	BL.	All claims for "Black Lung" benefits under Title 4, Part B, of the Federal Coal Mine He 923)	alth and Safety A	ct of 1969. (30 U.S.C.
863	DIWC	All claims filed by insured workers for disability insurance benefits under Title 2 of the all claims filed for child's insurance benefits based on disability. (42 U.S.C. 405 (g))	he Social Security	Act, as amended; plu
863	DIWW	All claims filed for widows or widowers insurance benefits based on disability unde amended. (42 U.S.C. 405 (g))	r Title 2 of the Soc	cial Security Act, as
864	SSID	All claims for supplemental security income payments based upon disability filed \boldsymbol{u} amended.	nder Title 16 of th	ne Social Security Act,
865	RSI	All claims for retirement (old age) and survivors benefits under Title 2 of the Social (42 U.S.C. 405 (g))	Security Act, as ar	mended.

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CIVIL COVER SHEET Page 3 of 3

UNITED STATES DISTRICT COURT CENTRAL DISTRICT OF CALIFORNIA

NOTICE OF ASSIGNMENT TO UNITED STATES JUDGES

This case has been assigned	to District Judge	Jesus G. Bernal	and to					
Magistrate Judge	David T. Bristow							
The case number on all doc	uments filed with the Cou	rt should read as follows:						
	5:14-cv-00667 JG	B-DTBx						
10.1. or or Cal. III is along. District Count for the Control District of								
Pursuant to General Order 05-07 of the United States District Court for the Central District of California, the assigned Magistrate Judge has been designated to hear discovery-related motions. All discovery-related motions should be noticed on the calendar of the Magistrate Judge.								
		Clerk, U. S. District Co	urt					
April 4, 2014 Date		By SBOURGEOIS Deputy Clerk						
ATTENTION								

A copy of this Notice must be served on all parties served with the Summons and Complaint (or, in cases removed from state court, on all parties served with the Notice of Removal) by the party who filed the Complaint (or Notice of Removal).